PA NT COOPERATION TREAT

From the INTERNATIONAL BUREAU

PCT	То:		
NOTIFICATION OF ELECTION (PCT Rule 61.2) Date of mailing (day/month/year)	Commissioner US Department of Commerce United States Patent and Trademark Office, PCT 2011 South Clark Place Room CP2/5C24 Arlington, VA 22202 ETATS-UNIS D'AMERIQUE in its capacity as elected Office		
26 February 2001 (26.02.01)			
International application No. PCT/FI00/00581	Applicant's or agent's file reference 50147		
International filing date (day/month/year) 28 June 2000 (28.06.00)	Priority date (day/month/year) 29 June 1999 (29.06.99)		
Applicant			
MARTIKAINEN, Mika	· ·		
1. The designated Office is hereby notified of its election made. X in the demand filed with the International Preliminary 24 January 20	y Examining Authority on: 001 (24.01.01)		
	Authorized officer		

Form PCT/IB/331 (July 1992)

Facsimile No.: (41-22) 740.14.35

The International Bureau of WIPO 34, chemin des Colombettes

1211 Geneva 20, Switzerland

R. E. Stoffel

Telephone No.: (41-22) 338.83.38

International application No. PCT/FI 00/00581

A. CLASSIFICATION OF SUBJECT MATTER

IPC7: C01G 51/04
According to International Patent Classification (IPC) or to both national classification and IPC B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: C01G

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

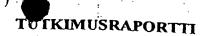
·	CONSIDERED TO BE RELEVANT	
Category*	Citation of document, with indication, where appropriate, of the relevant passages	
х	US 5569444 A (PHILIPPE DIANGUARE -	Relevant to claim N
	29 October 1996 (29.10.96), column 3, line 47 - line 61, claim 1, abstract	1-13
A	US 5057299 A (JEAN A. LOWN), 15 October 1991 (15.10.91), claim 1, abstract	1-13
		
1		

Further documents are listed in the continuation of Book Special categories of cited documents	ox C. X See patent family annex.
document defining the general state of the art which is not considered to be of particular relevance "E" criter document but published on or after the international filing date document which may throw doubts on priority claim(s) or which is special reason (as specified) document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed	"X" document of particular relevance: the claimed invention cannot be step when the document is taken alone "Y" document of particular relevance: the claimed invention cannot be step when the document is taken alone "Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is the combined with one or more other such documents, such combination of the particular in the second of the combination of the particular invention step when the document is the particular in the second of the particular invention of the parti
Date of the actual completion of the international search 24 October 2000 Name and mailing address of the ISA/ Swedish Patent Office Box 5055, S-102 42 STOCKHOLM	Date of mailing of the international search report 26 -10- 2000 Authorized officer

Moa Grönkvist/ELY

Telephone No. +46 8 782 25 00

PATENTTI- JA REKISTERIMALLITUS Patentti- ja innovaatiolinja



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TUTKITTU AINEISTO				
TOTRITTU AINEISTO	r.			• • • · · · · · · · · · · · · · · · · ·
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Kategoria* ⁾	Julkaisun tunnistetiedot		Koskee vaatimuksis
; X	US-A 5569444 (C 01 G 51/00)		1-13
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ja yksi t	tavuuden kannalta merkittävä julkaisu yksinään ta tavuuden kannalta merkittävä julkaisu, kun otetaa ii useampi-samaan kategoriaan kuuluva julkaisu ekniikan tasoa edustava julkaisu, ei kuitenkaan pal	n huomioon tämä	

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REC'D 08	OCT	2001	
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INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference	FOR FURTHER ACTION See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPE/		tion of Transmittal of International Examination Report (Form PCT/IPEA/416)			
50147/IR			Priority date (day/month/year)			
International application No.	International filing date (day)	monin/year)	•			
PCT/FI00/00581	28.06.2000		29.06.1999			
International Patent Classification (IPC) o	r national classification and IP	C ₇				
C 01 G 51/04						
Applicant Chamical C	ovet al					
OMG Kokkola Chemicals	oy ec ai					
This international preliminary example.	amination report has been prep	ared by this Intern	national Preliminary Examining			
This international preliminary example: Authority and is transmitted to the state of th	ne applicant according to Artic	le 36.	1			
2. This REPORT consists of a total	of 4 sheets, inc	cluding this cover	sheet.			
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has a smanded and are the	hasis for this report and/or site	es comaining icc	on, claims and/or drawings which have tifications made before this Authority			
(see Rule 70.16 and Section	on 607 of the Administrative In	structions under t	he PCT).			
These annexes consist of a total	of 1 sheets.					
This report contains indications in the second	relating to the following items:					
I Basis of the report	Basis of the report					
II Priority						
1 ! ! -	of opinion with regard to nove	ltv. inventive ster	and industrial applicability			
ļ ——	III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability					
V Reasoned statemen citations and explan	V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement					
VI Certain documents	cited		Ì			
VII Certain defects in t	he international application					
<u> </u>	ns on the international applicat	ion				
	T _r	Date of completion	of this report			
Date of submission of the demand		or completion				
24.01.2001		17.09.200	1			
		Authorized officer				
Name and mailing address of the IPEA	/56	aumorized officer				
Box 5055	1/9/8	Jan Carle	rud/ELY			
s-102 42 STOCKHOLM Facsimile No. 08-667 72 88		Telephone No. 08				
Form PCT/IPEA/409 (cover sheet) (Jan	Form PCT/IPEA/409 (cover sheet) (January 1998)					



INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No.
PCT/FI00/00581

I.	Basi	s of the report	
1.	With r	egard to the elements of the international application:*	Ì
		the international application as originally filed	
	\boxtimes	the description: , as originally fil	led
		pages 1-6 , filed with the dema	
		pages, filed with the letter of	
	- 7	pages, filed with the letter of	-
	\bowtie	the claims: pages , as originally fi	led
		, as amended (together with any statement) under article	: 19
		, 1110	
		pages 7, filed with the letter of 07.08.2001	-
	\boxtimes	the dequings:	1
	حع	, as originally fi	
		, filed with the defin	and
		pages, filed with the letter of	_
		the sequence listing part of the description: , as originally fi	iled
		pages	
		pages, filed with the letter of	
	the in Thes	regard to the language, all the elements marked above were available or furnished to this Authority in the language in we international application was filed, unless otherwise indicated under this item. e elements were available or furnished to this Authority in the following language	hich n is:
	in an	The amendments have resulted in the cancellation of: the description, pages the claims, Nos. the drawings, sheet/fig This report has been established as if (some of) the amendments had not been made, since they have been considered beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2 (c)).** placement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are refer this report as "originally filed" and are annexed to this report since they do not contain amendments (Rules 70.16 d 70.17). by replacement sheet containing such amendments must be referred to under item I and annexed to this report.	



INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No.
PCT/FI00/00581

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	Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability;
٧.	Reasoned statement duder Article 35(2) with 1 5g.
	citations and explanations supporting such statement

1. Statement

Novelty (N)	Claims Claims	1-9	YES NO
Inventive step (IS)	Claims Claims	1-9	YES NO
Industrial applicability (IA)	Claims Claims	1-9	YES NO

2. Citations and explanations (Rule 70.7)

Documents cited in the International Search Report:

- A) US 5569444 A
- B) US 5057299 A

This Report refers to the amended claims filed 7 August 2001.

The present invention relates to a hydroxide of cobalt or of cobalt and some other metal. It is intended to provide a product with high density and a large particle size.

Document A describes a hydroxide of cobalt, nickel and cadmium or zinc and its production. The main metal is nickel and the proportion of cobalt in the hydroxide is 1-8%. Ammonium ions are added to a solution of nitrates or sulphates of said metals in an amount such that the ratio of complexing agent to metal is approximately within the interval 0.5-3 specified in claim 1. The temperature is maintained between 80 and 85 degrees centigrade and the pH is regulated to a value of 9.2 +/- 0.1.

The present claim 1 does not specify the proportions of the metals in the hydroxide. However, it specifies the product as "cobaltous hydroxide or... ...of cobalt and some other metal". Thus, the invention is different from what is disclosed in A in that the main metal in the hydroxide is cobalt. Furthermore, the pH is 10-13 and the mixture is not heated.

Document B, which is cited in the description, is cited as a further example of prior art technique.

Therefore, the claimed invention is novel.



INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/FI00/00581

(To be used when the space in any of the preceding boxes is not sufficient)

Continuation of: V.

In view of what is disclosed in the cited documents, the cobaltous hydroxide cannot be considered to be obvious to a person skilled in the art. Therefore, the invention is considered to involve an inventive step. It is also considered to be industrially applicable.



Claims

- 1. Cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal, **characterised** in having a density of about 0.5-2.2 g/cm³, a particle size above about 1 μ m, typically about 1-20 μ m, and a specific surface of about 0.5-20 m²/g, and that it is prepared by adding a complexing agent and hydroxide ion under alkaline conditions to an aqueous cloride solution of cobalt or to an aqueous cloride solution of an alloy of cobalt and some other metal in order to form metal hydroxide, wherein the complexing agent is selected so as to form an ammonium complex with the metal ion, the molar ratio of complexing agent to metal being approx. 0.5-3 and the pH being adjusted in the range 10-13.
- 2. A cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal as defined in claim 1, **characterised** in that the pH is regulated in the range 11.2-12.0.
- 3. A cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal as defined in claim 1, **characterised** in that the other metal is nickel, manganese, magnesium or aluminium, or alloys of these.
- 4. A cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal as defined in claim 1, **characterised** in that the concentration of the cloride solution is in the range from 10 to 120 g/l calculated on the total metal content.
- 5. A cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal as defined in claim 1, **characterised** in that the complexing agent is ammonium sulphate or aqueous ammonia
- 6. A cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal as defined in claim 1, **characterised** in that the molar ratio of complexing agent to metal is approx. 1.5-2.
- 7. A cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal as defined in claim 1, **characterised** in that NaOH is used for pH regulation.
- 8. A cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal as defined in claim 1, **characterised** in that the reaction is carried out at a temperature of about 40-90 °C.
- 9. A cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal as defined in claim 8, **characterised** in that the reaction is carried out at a temperature of about 70 °C.



From the INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

То:
Berggren Oy Ab P.O. Box 16 FIN-00101 Helsinki
Finland Berggren Oy No
19-06-2001
\v'

PCT

WRITTEN OPINION

(PCT Rule 66)

, w/ r	2001			
(V)		Date of mailing (day/month/year)	1 5 -06- 2001	
Applicant's or agent's file reference 50147/IR		REPLY DUE	within 60 days from the above date of mailing	14.8.
International application No. International filing date PCT/FI00/00581 28.06.2000		(day/month/year)	Priority date (day/month/year) 29.06.1999	
International Patent Classification (IP	C) or both national classifica	tion and IPC7		
C 01 G 51/04		<u> </u>		
Applicant				
OMG Kokkola Chemica	ls Oy et al			
				

1.	This written opinion is the first (first, etc.) drawn by this International Preliminary Examining Authority.			
2.	2. This opinion contains indications relating to the following items:			
	Basis of the report			
	II Priority			
	III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability			
	IV Lack of unity of invention			
	Neasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement			
	VI Certain documents cited			
	VII Certain defects in the international application			
	VIII Certain observations on the international application			
3.	The applicant is hereby invited to reply to this opinion.			
	When? See the time limit indicated above. The applicant may, before the expiration of that time limit, request this Author to grant an extension, see Rule 66.2(d).	ıty		
	How? By submitting a written reply, accompanied, where appropriate, by amendments, according to Rule 66.3. For the form and the language of the amendments, see Rules 66.8 and 66.9.			
	Also For an additional opportunity to submit amendments, see Rule 66.4. For the examiner's obligation to consider amendments and/or arguments, see Rule 66.4bis. For an informal communication with the examiner, see Rule 66.6.			
If no reply is filed, the international preliminary examination report will be established on the basis of this opinion.				
4.	4. The final date by which the international preliminary examination report must be established according to Rule 69.2 is: 29.10.2001			

Name and mailing address of the IPEA/SE		Authorized officer
Patent- och registreringsverket	Telex	
Box 5055	17978	a lama /mrv
S-102 42 STOCKHOLM	PATOREG-S	Jan Carlerud/ELY
Facsimile No. 08-667 72 88		Telephone No. 08-782 25 00



Intern	al application No.
PCT/	FI00/00581

With regard to the elements of the international application:* With regard to the elements of the international application as originally filed	I.		sis of the opinion	
the description: pages	1.	With		
pages filed with the letter of filed with the letter of filed with the demand pages filed with the letter of filed with the demand pages filed with the letter of filed with the letter of filed with the demand pages filed with the letter of filed with the demand pages filed with the letter of filed with the demand pages filed with the letter of filed with the letter of filed with the demand pages filed with the letter of filed with the demand pages filed with the letter of filed		\boxtimes	the international application as originally filed	
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the claims: pages				
the claims: pages			pagesfiled	with the letter of
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pages			pages , as am	ended (together with any statement) under article 19
the drawings:				,
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the sequence listing part of the description: pages pages pages , filed with the letter of 2. With regard to the language, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item. These elements were available or furnished to this Authority in the following language English which is: the language of a translation furnished for the purposes of international search (under Rule 23.1(b)). the language of publication of the international application (under Rule 48.3(b)). the language of the translation furnished for the purposes of international preliminary examination (under Rules 55.2 and/or 55.3). 3. With regard to any nucleotide and/or amino acid sequence disclosed in the international application, the written opinion was drawn on the basis of the sequence listing: contained in the international application in printed form. filed together with the international application in computer readable form. furnished subsequently to this Authority in computer readable form. furnished subsequently to this Authority in computer readable form. The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished. The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished. 4. The amendments have resulted in the cancellation of: the description, pages the claims, Nos. the drawings, sheet/fig This opinion has been drawn as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2 (c)).			pages, filed	with the letter of
pages				
2. With regard to the language, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item. These elements were available or furnished to this Authority in the following language			pages	
2. With regard to the language, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item. These elements were available or furnished to this Authority in the following language			pages	, illed with the demand
the international application was filed, unless otherwise findeated unter a terms were available or furnished to this Authority in the following language English which is: the language of a translation furnished for the purposes of international search (under Rule 23.1(b)). the language of publication of the international application (under Rule 48.3(b)). the language of the translation furnished for the purposes of international preliminary examination (under Rules 55.2 and/ or 55.3). 3. With regard to any nucleotide and/or amino acid sequence disclosed in the international application, the written opinion was drawn on the basis of the sequence listing: contained in the international application in printed form. filed together with the international application in computer readable form. furnished subsequently to this Authority in written form. furnished subsequently to this Authority in computer readable form. The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished. The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished. 4. The amendments have resulted in the cancellation of: the description, pages	l		pages, filed	with the letter of
the claims, Nos. the drawings, sheet/fig This opinion has been drawn as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2 (c)). * Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to		the in Thes	the language of a translation furnished for the purposes of international the language of publication of the international application (under Rul the language of the translation furnished for the purposes of internation or 55.3). ith regard to any nucleotide and/or amino acid sequence disclosed in the awn on the basis of the sequence listing: contained in the international application in printed form. filed together with the international application in computer readable furnished subsequently to this Authority in written form. The statement that the subsequently furnished written sequence listing international application as filed has been furnished. The statement that the information recorded in computer readable for been furnished.	clanguage English which is: al search (under Rule 23.1(b)). the 48.3(b)). and preliminary examination (under Rules 55.2 and/ international application, the written opinion was form. g does not go beyond the disclosure in the
		* Re	the description, pages the claims, Nos. the drawings, sheet/fig This opinion has been drawn as if (some of) the amendments had not beyond the disclosure as filed, as indicated in the Supplemental Box Replacement sheets which have been furnished to the receiving Office in re-	(Ruic 70.2 (0)).



International application No.
PCT/FI00/00581

		:
v -	/. Reasoned statement under Rule 66.2(a)(ii) with regard to novel	ty, inventive step of industrial approximation,
٠.	citations and explanations supporting such statement	

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Citations and explanations

Documents cited in the International Search Report:

- A) US 5569444 A
- B) US 5057299 A

The present invention relates to a method for preparing a hydroxide of cobalt or of cobalt and some other metal with high density and a large particle size and to the product produced with the method.

Document A describes a method for producing hydroxide of cobalt, nickel and cadmium or zinc. Ammonium ions are added to a solution of nitrates or sulphates of said metals in an amount such that the ratio of complexing agent to metal is approximately within the interval 0.5-3 specified in claim 1. The pH is regulated to a value of 9.2 + - 0.1.

Thus, the claimed method is different from what is disclosed in A in that the pH is 10-13. It is, however, considered to be an obvious measure to a person skilled in the art to perform experiments to find out what effect changes to parameters such as pH, temperature or concentrations have to the product. Therefore, the method of claims 1-10 is not considered to involve an inventive step.

The product specified in claims 11-13 does not differ from the product disclosed in A, see column 3, lines 47-61. Therefore, the invention of claims 11-13 lacks novelty.

Document B, which is cited in the description, is cited as a further example of prior art technique.



Internation No.
PCT/FI00/00581

VII. Certain defects in the international application

The following defects in the form or contents of the international application have been noted:

Claim 13 refers to claims 11 and 12, which both are independent claims. This is not allowed. A dependent claim can refer only to one independent claim.



Managing Director 0/018986

531 Rec'd PCT/PTC 19 DEC 200 POYLLISYYSMALLIT - PATENTS 7 August 2001

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L. Nordin

· PATENTIT

J. Kupiainen M. Brax E. Heikkinen T. Laako B. Lassenius T. Pelin I. Risku O-P. Saiionmaa J. Svensson B. Träskman J. Jordnen M Karttunen S. Kuisma M. Laajalanti V. Tognetty S. Ylätalo

> MALLIT: - DESIGNS

N. Mikander L. Valjakka

· TAVARAMERKIT LAKIASIAT · TRADEMARKS.

E-M. Söderström** S. Aspoia

LEGAL MATTERS: P. Kolve* H. Halmetoja** S. Henn** 1. Karlsson

Patent- och registreringsverket Box 5055 S-102 42 Stockholm Sweden

FAX: +46-8-667 7288 (5 pages) Confirmation by mail

Authorized Officer: Jan Carlerud/ELY

Our ref: 50147/IR/MG

REPLY TO WRITTEN OPINION INTERNATIONAL PATENT APPLICATION PCT/F100/00581 APPLICANT: OMG KOKKOLA CHEMICALS OY

On account of the Written Opinion issued on 15 June 2001 we submit the following:

We enclose a new claim set directed to the cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal, i.e. to the product itself. The aqueous brine has been amended to a chloride solution as illustrated in the examples.

In applications of the accumulator industry the capacity of the accumulators can be raised by using cobaltous hydroxide with maximum density as an additive in accumulators. In addition it is important that the cobaltous hydroxide used has a small specific surface and a high density. The cobaltous hydroxide of claim 1 has both and also large particle size.

The conventional cobaltous hydroxide is very fine-grained and because of this small particle size it also has large spesific surface.

The larger particle size, and specifically the possibility for adjusting the particle size, is achieved by choice of the complexing agent and process parameters, as in claim 1.

In the method of US patent 5,057,299 the mixture is heated and kept in hydrothermal conditions until the cobaltous hydroxide precipitates. With the method of this US patent, the particle size increases at higher temperatures. The method of the patent allows the particle size to be controlled only in the range from approx. 0.05 to $0.5~\mu m$ by means of the temperature. Temperature is not critical parameter in the present invention.

Berggren Oy Ab Oscite • Address: PL 16 • P.O.Box 16

FIN-00101 Helsinki FINLAND *European Patent Attorney *European Trademark Attorney Käyntiosoite • Office: Graniittitalo Jaakonkatu 3 A Helsinki

6 Nat. (09) 693 701 Int. +358 9 693 701 Fax +358 9 693 3944 email.box@berggren.fi http://www.berggren.fi Pankit • Bankers: NORDEA 157330-15411 MRITF!HH SWIFT 800017-90104 SAMPO PSPBFIHH. SWIFT

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Kotipaikka

Helsinki

Berggren Y

In the US patent 5,569,444 a method for preparing **nickel** hydroxide is presented. There may be only 8 % cobalt among the nickel. Further solutions of nitrates or sulphates are used in the US patent, whereas in the present invention chloride solution is used. Also the pH is lower in the US patent than in the present invention.

The cobaltous hydroxide of claim 1 has considerably larger particle size than the metal hydroxides of the cited prior art, and this is achieved by choice of the complexing agent and process parameters, as in claim 1.

BERGGREN OY AB

Ira Risku

Patent Agent

Encls

a set of new claims 1 to 9 in triplicate

531 Rec'd PCTA

19 DEC 2001

Claims

- 1. Cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal, characterised in having a density of about 0.5-2.2 g/cm³, a particle size above about 1 µm, typically about 1-20 µm, and a specific surface of about 0.5-20 m²/g, and that it is prepared by adding a complexing agent and hydroxide ion under alkaline conditions to an aqueous cloride solution of cobalt or to an aqueous cloride solution of an alloy of cobalt and some other metal in order to form metal hydroxide, wherein the complexing agent is selected so as to form an ammonium complex with the metal ion, the molar ratio of complexing agent to metal being approx. 0.5-3 and the pH being adjusted in the range 10-13.
- 2. A cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal as defined in claim 1, characterised in that the pH is regulated in the range 11.2-12.0.
- 3. A cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal as defined in claim 1, **characterised** in that the other metal is nickel, manganese, magnesium or aluminium, or alloys of these.
- 4. A cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal as defined in claim 1, characterised in that the concentration of the cloride solution is in the range from 10 to 120 g/l calculated on the total metal content.
- 5. A cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal as defined in claim 1, characterised in that the complexing agent is ammonium sulphate or aqueous ammonia
- 6. A cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal as defined in claim 1, characterised in that the molar ratio of complexing agent to metal is approx. 1.5-2.
- 7. A cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal as defined in claim 1, characterised in that NaOH is used for pH regulation.
- 8. A cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal as defined in claim 1, **characterised** in that the reaction is carried out at a temperature of about 40-90 °C.
- 9. A cobaltous hydroxide or alloy hydroxide formed of cobalt and some other metal as defined in claim 8, **characterised** in that the reaction is carried out at a temperature of about 70 °C.

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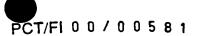
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PCT REQUEST

Original (for SUBMISSION) - printed on 28.06.2000 09:26:26 AM

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0	For receiving Office use only	PCT/FI 0 0 / 0 0 5 8 1		
0-1	International Application No.	PC///100/80581		
0-2	International Filing Date	2 8 JUN 2000 (2 8 -06- 2000)		
0-3	Name of receiving Office and "PCT International Application"	The Finnish Patent Office PCT International Application		
0-4	Form - PCT/RO/101 PCT Request			
0-4-1	Prepared using	PCT-EASY Version 2.90 (updated 10.05.2000)		
0-5	Petition The undersigned requests that the present international application be processed according to the Patent Cooperation Treaty			
0-6	Receiving Office (specified by the applicant)	National Board of Patents and Registration (Finland) (RO/FI)		
0-7	Applicant's or agent's file reference	50147		
		PROCESS FOR MAKING HIGH DENSITY AND LARGE PARTICLE SIZE COBALT HYDROXIDE OR COBALT MIXED HYDROXIDES AND A PRODUCT MADE BY THIS PROCESS		
	Applicant			
II-1	This person is:	applicant only		
II-2	Applicant for	all designated States except US		
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II-6	State of nationality	FI		
o II-7	State of residence	FI		
III-1	Applicant and/or inventor			
III-1-1	This person is:	applicant and inventor		
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III-1-5	Address:	Hakalahdenkatu 83 B 22 FIN-67100 Kokkola		
		Finland		
III-1-6	State of nationality	FI		



PCT REQUEST

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IV-1	Agent or common representative; or address for correspondence The person identified below is hereby/has been appointed to act on behalf of the applicant(s) before the competent International Authorities as:	agent
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V	Designation of States	
V-1	Regional Patent (other kinds of protection or treatment, if any, are specified between parentheses after the designation(s) concerned)	AP: GH GM KE LS MW MZ SD SL SZ TZ UG ZW and any other State which is a Contracting State of the Harare Protocol and of the PCT
		EA: AM AZ BY KG KZ MD RU TJ TM and any
		other State which is a Contracting State
		of the Eurasian Patent Convention and of
		the PCT
		EP: AT BE CH&LI CY DE DK ES FI FR GB GR
		IE IT LU MC NL PT SE and any other State
		which is a Contracting State of the
		European Patent Convention and of the
		OA: BF BJ CF CG CI CM GA GN GW ML MR NE
		SN TD TG and any other State which is a
		member State of OAPI and a Contracting
		State of the PCT
V-2	National Patent	AE AG AL AM AT AU AZ BA BB BG BR BY BZ
	(other kinds of protection or treatment, if any, are specified between parentheses	CA CH&LI CN CR CU CZ DE DK DM DZ EE ES
	after the designation(s) concerned)	FI GB GD GE GH GM HR HU ID IL IN IS JP
		KE KG KP KR KZ LC LK LR LS LT LU LV MA
	1	MD MG MK MN MW MX MZ NO NZ PL PT RO RU
	· ·	SD SE SG SI SK SL TJ TM TR TT TZ UA UG
V-5	Precautionary Designation Statement	US UZ VN YU ZA ZW
V-5	In addition to the designations made under items V-1, V-2 and V-3, the applicant also makes under Rule 4.9(b)	
	all designations which would be	
	permitted under the PCT except any designation(s) of the State(s) indicated	
	under item V-6 below. The applicant	
	declares that these additional	
	declares that those additional designations are subject to confirmation	
	designations are subject to confirmation and that any designation which is not	
	designations are subject to confirmation and that any designation which is not confirmed before the expiration of 15 months from the priority date is to be	
	designations are subject to confirmation and that any designation which is not confirmed before the expiration of 15	

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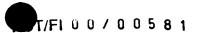
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VI-1	Priority claim of earlier national		
VI-1-1	application Filing date	29 June 1999 (29.06.	1000)
VI-1-2	Number	991478	1999)
VI-1-3	Country		
VI-1-3	<u> </u>	FI	
	Priority document request The receiving Office is requested to prepare and transmit to the International Bureau a certified copy of the earlier application(s) identified above as item(s):	VI-1	
VII-1	International Searching Authority Chosen	Swedish Patent Offic	e (ISA/SE)
VIII	Check list	number of sheets	electronic file(s) attached
VIII-1	Request	4	-
VIII-2	Description	6	-
VIII-3	Claims	2	_
VIII-4	Abstract	1	50147.txt
VIII-5	Drawings	1	-
VIII-7	TOTAL	14	· · · · · · · · · · · · · · · · · · ·
	Accompanying items	paper document(s) attached	electronic file(s) attached
VIII-8	Fee calculation sheet	√	-
VIII-9	Separate signed power of attorney	✓	_
VIII-10	Copy of general power of attorney	√	_
VIII-16	PCT-EASY diskette	_	diskette
VIII-17	Other (specified):	Copy of Official	-
		Action in FI 991478	
VIII-18	Figure of the drawings which should accompany the abstract		
VIII-19	application	Finnish	
IX-1	Signature of applicant or agent	In Est	
IX-1-1	Name	BERGGREN OY AB	
IX-1-2	Name of signatory	Ira Risku	
IX-1-3	Capacity	Patent Attorney	

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10-1	Date of actual receipt of the purported international application	2 8	3 JUN 2000	(2 g -06- 2000)	
10-2	Drawings:		· · · · · · · · · · · · · · · · · · ·		
10-2-1	Received				
10-2-2	Not received				
10-3	Corrected date of actual receipt due to later but timely received papers or drawings completing the purported international application				
10-4	Date of timely receipt of the required corrections under PCT Article 11(2)				
10-5	International Searching Authority	ISA/SE			
10-6	Transmittal of search copy delayed until search fee is paid		<u>, , , , , , , , , , , , , , , , , , , </u>		= .



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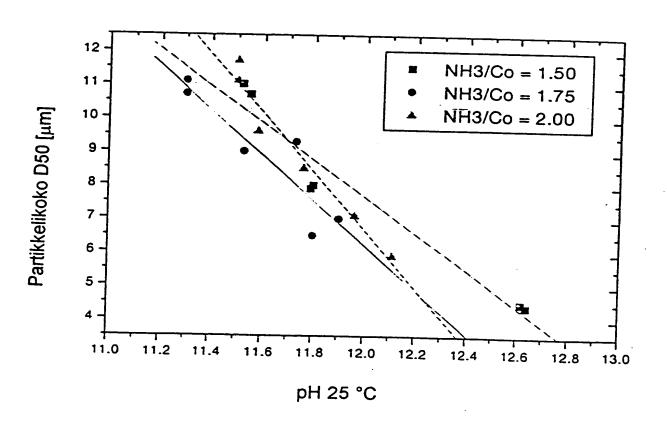
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11-1	Date of receipt of the record copy by	2.5	11111	9000	1 2 5, 67, 60)
	the International Bureau	25	<u> JULY</u>	2000	\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \



Kuva 1

Menetelmä korkean tilavuuspainon ja suuren partikkelikoon omaavan kobolttihydroksidin tai kobolttimetalliseoshydroksidin valmistamiseksi ja menetelmällä saatu tuote

Keksintö koskee menetelmää korkean tilavuuspainon ja suuren partikkelikoon omaavan kobolttihydroksidin tai koboltin ja jonkin muun metallin seoshydroksidin valmistamiseksi. Keksintö koskee myös tällä menetelmällä aikaansaatua tuotetta.

Kobolttihydroksidia käytetään monissa sovellutuksissa, esimerkiksi elektroniikkateollisuudessa lisäaineena ladattavissa NiMH- ja NiCd-akuissa. Lisäksi se soveltuu käytettäväksi oksidituotteiden, kuten LiCoO₂ ja LiCoMO₂, valmistuksessa prekursorina (M tarkoittaa metallia). Kobolttihydroksidia käytetään myös katalyyttinä tai prekursorina katalyyttituotannossa. Edellä mainittuihin tarkoituksiin soveltuu käytettäväksi myöskin hydroksidituote, jossa koboltin ohella on jotain muuta metallia, kuten nikkeliä, mangaania, mangnesiumia tai alumiinia.

Alalla tunnetaan monia menetelmiä kobolttihydroksidin valmistamiseksi. US-patentin 5 057 299 mukaisessa menetelmässä kobolttihydroksidia valmistetaan yhdistämällä koboltti-ioni kompleksoivan aineen kanssa, jolloin muodostuu vesiliukoinen kobolttikompleksi. Tähän lisätään hydroksidi-ionia ja seosta kuumennetaan ja pidetään hydrotermaalisissa olosuhteissa, kunnes kobolttihydroksidi saostuu. Tämän US-patentin menetelmällä partikkelikoko saadaan kasvamaan korkeampia lämpötiloja käytettäessä. Partikkelikokoa voidaan patentin mukaisella menetelmällä lämpötilan avulla säätää kuitenkin vain välillä noin 0,05-0,5 μm.

Akkuteollisuuden sovellutuksissa NiCd- ja NIMH-akuilta vaaditaan hyvää kapasiteettia. Kapasiteettia voidaan nostaa käyttämällä akkujen lisäaineena kobolttihydroksidia, jonka tilavuuspaino on mahdollisimman korkea. Lisäksi oksidituotteiden valmistuksessa on tärkeää, että käytetyllä kobolttihydroksilla on alhainen ominaispinta-ala ja korkea tilavuuspaino. Esillä olevan keksinnön tarkoituksena on saada aikaan menetelmä, jolla voidaan valmistaa suuren partikkelikoon omaavia kobolttihydroksidi-partikkeleita tai koboltin ja jonkin muun metallin seoshydroksidipartikkeleita helposti kontrolloitavalla menetelmällä. Tarkoituksena on saada aikaan kobolttihydroksidipartikkeleita tai koboltin ja jonkin muun metallin seoshydroksidipartikkeleita, joiden partikkelikoko on yli 1 μm, edullisesti yli 3 μm. Tämä on keksinnön mukaisesti saatu aikaan siten kuin on esitetty oheisissa patenttivaatimuksissa.

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Keksinnön mukaisella menetelmällä voidaan valmistaa paitsi kobolttihydroksidia myös koboltin ja jonkin muun metallin tai metalleiden seoshydroksidia. Muina metalleina voidaan käyttää esimerkiksi nikkeliä, mangaania, mangnesiumia tai alumiinia tai näiden seoksia käyttötarkoituksen mukaan.

Keksinnön mukaisessa menetelmässä käytetään lähtöaineena koboltin tai koboltin ja 5 seosmetallien vesipitoisia suolaliuoksia sulfaatin, nitraatin tai kloridin muodossa. Liuoksen konsentraatio voi vaihdella välillä 10-120 g/l kokonaismetallipitoisuuden suhteen laskettuna.

Tähän metallisuolaliuokseen lisätään sellaista kompleksinmuodostaja-ainetta, joka 10 muodostaa ammoniumkompleksin metalli-ionin kanssa. Kompleksinmuodostajaaine voi olla ammoniumsulfaattia, vesipitoista ammoniakkia tai jokin muu ammoniumionilähde. Periaatteessa kompleksinmuodostaja-aine voi olla myös jotain muuta metallin kanssa kompleksin muodostavaa ainetta, kuten esimerkiksi EDTA:ta. Esillä olevassa keksinnössä ammoniumioni on kuitenkin havaittu hyväksi kompleksin-15 muodostaja-aineeksi sen halpuuden ja hyvien kompleksinmuodostusominaisuuksiensa takia. Reaktoriin syötettävän ammoniumkompleksinmuodostaja-aineen ja metallin välinen moolisuhde on edullisesti noin 0,5-3, edullisimmin noin 1,5-2,0. Reaktio suoritetaan alkalisissa olosuhteissa ja pH:n säätämiseen käytetään edullisesti NaOH:ia. Edullinen pH-arvo on välillä 10-13, edullisimmillaan pH on välillä 11,2-12,0. Reaktio suoritetaan lämpötilassa noin 40-90 °C, edullisesti lämpötilassa noin 70 °C. Keksinnön mukaisella menetelmällä saadaan kobolttihydroksidia, jonka tilavuuspaino on noin 0,5-2,2 g/cm³, partikkelikoko yli noin 1 μm, tyypillisesti noin 1-20 μ m ja ominaispinta-ala noin 0,5-20 m²/g.

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Keksinnön mukaisella menetelmällä saadut partikkelit ovat muodoltaan heksagonaalisia levymäisiä kobolttihydroksidipartikkeleita. Keksinnön yhteydessä havaittiin, 25 että kloridiliuoksilla saadaan paksumpia hiukkasia kuin sulfaattipohjaisilla liuoksilla, kun muut reaktio-olosuhteet pidetään samoina. Paksumpien hiukkasten katsotaan soveltuvan paremmin esimerkiksi litiumkobolttioksidin valmistukseen.

Keksinnön mukaisessa menetelmässä kobolttihydroksidin partikkelikokoa voidaan säätää ammoniumionin ja koboltti-ionin moolisuhteen avulla ja käytetyn pH:n avul-30 la. Kuvassa 1 on esitetty partikkelikoon muutos pH:n funktiona erilaisilla NH₃ioni/Co-suhteilla. Kuvasta nähdään, että pH:n nostaminen pienentää partikkelikokoa, ja että mitä korkeampi on NH3-ioni/Co-suhde sitä merkittävämpi vaikutus pH:lla on partikkelikokoon.

Seuraavassa keksintöä on valaistu esimerkkien avulla. Esimerkeissä 1-3 on menetelmä suoritettu vertailuna ilman kompleksinmuodostusta ammoniumionin kanssa. Esimerkeissä 4-6 on kuvattu keksinnön mukainen menetelmä, jossa muodostetaan ammoniumkompleksi koboltin kanssa. Esimerkissä 7 on kuvattu keksinnön mukainen menetelmä, jossa on käytetty koboltin lisäksi nikkeliä. Esimerkeissä on reaktiolämpötilana käytetty 70 °C.

Vertailuesimerkki 1

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CoCl₂:n vesiliuosta (30 g/l Co) syötettiin jatkuvasti ensimmäiseen reaktoriin. pH pidettiin vakioarvossa 11,8-12,0 lisäämällä natriumhydroksidiliuosta. Ensimmäisen reaktorin ylivirtaus johdettiin toiseen reaktoriin, jossa virtaukseen lisättiin natriumhydroksidia siten, että pH oli 13,5. Toisen reaktorin ylivirtaus suodatettiin ja pestiin vedellä. Pesty Co(OH)₂-kakku kuivattiin.

Kuivatun Co(OH)₂:n keskimääräinen partikkelikoko D50 oli 0,9 μm mitattuna Malvern Mastersizer -partikkelikokoanalysaattorilla (mittaus suoritettiin laserdiffraktiolla vesipitoisesta Co(OH)₂-lietteestä). Tilavuuspaino oli 0,5 g/cm³ (ASTM B527-93). Ominaispinta-ala (BET ASTM D4567-86) oli 30 m²/g.

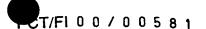
Vertailuesimerkki 2

CoCl₂:n vesiliuosta (30 g/l Co) syötettiin jatkuvasti ensimmäiseen reaktoriin. pH pidettiin vakioarvossa 11,6-11,8 lisäämällä natriumhydroksidiliuosta. Ensimmäisen reaktorin ylivirtaus johdettiin toiseen reaktoriin, jossa virtaukseen lisättiin natriumhydroksidia siten, että pH oli 13,5. Toisen reaktorin ylivirtaus suodatettiin ja pestiin vedellä. Pesty Co(OH)₂-kakku kuivattiin.

Kuivatun Co(OH)₂:n keskimääräinen partikkelikoko D50 oli 1,1 μm mitattuna Malvern Mastersizer -partikkelikokoanalysaattorilla (mittaus suoritettiin laserdiffraktiolla vesipitoisesta Co(OH)₂-lietteestä). Tilavuuspaino oli 0,7 g/cm³ (ASTM B527-93). Ominaispinta-ala (BET ASTM D4567-86) oli 29 m²/g.

Vertailuesimerkki 3

CoCl₂:n vesiliuosta (30 g/l Co) syötettiin jatkuvasti ensimmäiseen reaktoriin. pH pidettiin vakioarvossa 11,2-11,4 lisäämällä natriumhydroksidiliuosta. Ensimmäisen reaktorin ylivirtaus johdettiin toiseen reaktoriin, jossa virtaukseen lisättiin natriumhydroksidia siten, että pH oli 13,5. Toisen reaktorin ylivirtaus suodatettiin ja pestiin vedellä. Pesty Co(OH)₂-kakku kuivattiin.



Kuivatun Co(OH)₂:n keskimääräinen partikkelikoko D50 oli 1,9 μm mitattuna Malvern Mastersizer -partikkelikokoanalysaattorilla (mittaus suoritettiin laserdiffraktiolla vesipitoisesta Co(OH)₂-lietteestä). Tilavuuspaino oli 0,6 g/cm³ (ASTM B527-93). Ominaispinta-ala (BET ASTM D4567-86) oli 35 m²/g.

5 Esimerkki 4

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CoCl₂:n vesiliuosta (30 g/l Co) syötettiin jatkuvasti ensimmäiseen reaktoriin, jossa oli ammoniakin vesiliuosta siten, että metallin moolisuhde ammoniumiin oli 2. pH pidettiin vakioarvossa 11,8-12,0 lisäämällä natriumhydroksidiliuosta. Ensimmäisen reaktorin ylivirtaus johdettiin toiseen reaktoriin, jossa virtaukseen lisättiin natriumhydroksidia siten, että pH oli 13,5. Toisen reaktorin ylivirtaus suodatettiin ja pestiin vedellä. Pesty Co(OH)₂-kakku kuivattiin.

Kuivatun Co(OH)₂:n keskimääräinen partikkelikoko D50 oli 1,8 μm mitattuna Malvern Mastersizer -partikkelikokoanalysaattorilla (mittaus suoritettiin laserdiffraktiolla vesipitoisesta Co(OH)₂-lietteestä). Tilavuuspaino oli 0,7 g/cm³ (ASTM B527-93). Ominaispinta-ala (BET ASTM D4567-86) oli 5,8 m²/g.

Esimerkki 5

CoCl₂:n vesiliuosta (30 g/l Co) syötettiin jatkuvasti ensimmäiseen reaktoriin, jossa oli ammoniakin vesiliuosta siten, että metallin moolisuhde ammoniumiin oli 2. pH pidettiin vakioarvossa 11,6-11,8 lisäämällä natriumhydroksidiliuosta. Ensimmäisen reaktorin ylivirtaus johdettiin toiseen reaktoriin, jossa virtaukseen lisättiin natriumhydroksidia siten, että pH oli 13,5. Toisen reaktorin ylivirtaus suodatettiin ja pestiin vedellä. Pesty Co(OH)₂-kakku kuivattiin.

Kuivatun Co(OH)₂:n keskimääräinen partikkelikoko D50 oli 3,9 μm mitattuna Malvern Mastersizer -partikkelikokoanalysaattorilla (mittaus suoritettiin laserdiffraktiolla vesipitoisesta Co(OH)₂-lietteestä). Tilavuuspaino oli 1,2 g/cm³ (ASTM B527-93). Ominaispinta-ala (BET ASTM D4567-86) oli 2,6 m²/g.

Esimerkki 6

CoCl₂:n vesiliuosta (30 g/l Co) syötettiin jatkuvasti ensimmäiseen reaktoriin, jossa oli ammoniakin vesiliuosta siten, että metallin moolisuhde ammoniumiin oli 2. pH pidettiin vakioarvossa 11,2-11,4 lisäämällä natriumhydroksidiliuosta. Ensimmäisen reaktorin ylivirtaus johdettiin toiseen reaktoriin, jossa virtaukseen lisättiin natrium-

hydroksidia siten, että pH oli 13,5. Toisen reaktorin ylivirtaus suodatettiin ja pestiin vedellä. Pesty Co(OH)₂-kakku kuivattiin.

Kuivatun Co(OH)₂:n keskimääräinen partikkelikoko D50 oli 7,4 µm mitattuna Malvern Mastersizer -partikkelikokoanalysaattorilla (mittaus suoritettiin laserdiffraktiolla vesipitoisesta Co(OH)₂-lietteestä). Tilavuuspaino oli 1,7 g/cm³ (ASTM B527-93). Ominaispinta-ala (BET ASTM D4567-86) oli 1,8 m²/g.

Esimerkki 7

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Valmistettiin seosmetallisuolaliuos sisältäen CoCl₂- ja NiCl₂-liuoksia (30 g/l Co ja 8 g/l Ni). Liuosta syötettiin jatkuvasti ensimmäiseen reaktoriin, jossa oli ammoniakin vesiliuosta siten, että metallin moolisuhde ammoniumiin oli 2. pH pidettiin vakioarvossa 11,2-11,4 lisäämällä natriumhydroksidiliuosta. Ensimmäisen reaktorin ylivirtaus johdettiin toiseen reaktoriin, jossa virtaukseen lisättiin natriumhydroksidia siten, että pH oli 13,5. Toisen reaktorin ylivirtaus suodatettiin ja pestiin vedellä. Pesty kakku kuivattiin.

- Kemiallinen analyysi osoitti, että seosmetallihydroksidia oli saostunut. Kuivatun Co0,8Ni0,2(OH)₂:n keskimääräinen partikkelikoko D50 oli 6,9 μm mitattuna Malvern Mastersizer -partikkelikokoanalysaattorilla (mittaus suoritettiin laserdiffraktiolla vesipitoisesta lietteestä). Tilavuuspaino oli 1,6 g/cm³ (ASTM B527-93). Ominaispinta-ala (BET ASTM D4567-86) oli 3,2 m²/g.
- Vertailun helpottamiseksi alla on esitetty taulukon muodossa edellä kuvattujen esimerkkien tuotteiden fysikaaliset ominaisuudet. Taulukossa on lisäksi esitetty tuotteiden kidekoko mitattuna röntgendiffraktiometrillä.

Taulukko 1

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Esimerkki	Saostus- pH	Kompleksin- muodostus- aine	Keskimää- räinen partikkeli- koko (µm)	Tilavuus- paino (g/cm³)	Ominais- pinta-ala (m²/g)		XRD (101) (nm)
Vertailu- esimerkki 1	11,8-12,0	ei mitään	0,9	0,5	30	23	27
Vertailu- esimerkki 2	11,6-11,8	ei mitään	1,1	0,7	29	26	30
Vertailu- esimerkki 3	11,2-11,4	ei mitään	1,9	0,6	35	27	32
Esimerkki 4	11,8-12,0	NH ₃ -ioni	1,8	0,7	5,8	50	49
Esimerkki 5	11,6-11,8	NH₃-ioni	3,9	1,2	2,6	56	56
Esimerkki 6	11,2-11,4	NH ₃ -ioni	7,4	1,7	1,8	61	64
Esimerkki 7	11,2-11,4	NH ₃ -ioni	6,9	1,6	3,2	59	57

Taulukosta voidaan nähdä, että käytettäessä kompleksinmuodostaja-aineena ammoniumionia, saatiin tuotteelle suurempi partikkelikoko ja tilavuuspaino kuin ilman kompleksinmuodostajaa. Poikkeuksena oli esimerkin 4 tuote, jonka partikkelikoko ja tilavuuspaino vastasivat suunnilleen vertailuesimerkkien tuotteiden partikkelikokoa ja tilavuuspainoa. Tosin tälläkin tuotteella ominaispinta-ala ja kidekoko vastasivat esimerkkien 5-7 tuotteiden arvoja.

Edellä on esitetty eräitä keksinnön sovelluksia. Keksintöä luonnollisesti ei rajoiteta edellä esitettyihin esimerkkeihin, vaan keksinnön mukaista periaatetta voidaan muunnella patenttivaatimusten suoja-alan puitteissa.

Patenttivaatimukset

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- 1. Menetelmä korkean tilavuuspainon ja suuren partikkelikoon omaavan kobolttihydroksidin tai koboltin ja jonkin muun metallin seoshydroksidin valmistamiseksi, jossa koboltin vesipitoiseen suolaliuokseen tai koboltin ja jonkin muun metallin seoksen vesipitoiseen suolaliuokseen lisätään alkalisissa olosuhteissa kompleksinmuodostaja-ainetta ja hydroksidi-ionia metallihydroksidin muodostamiseksi, tunnettu siitä, että kompleksinmuodostaja-aine valitaan siten, että se muodostaa ammoniumkompleksin metalli-ionin kanssa, kompleksinmuodostaja-aineen ja metallin välinen moolisuhde on noin 0,5-3, ja pH säädetään välille 10-13.
- 2. Patenttivaatimuksen 1 mukainen menetelmä, tunnettu siitä, että pH säädetään välille 11,2-12,0.
 - 3. Patenttivaatimuksen 1 mukainen menetelmä, tunnettu siitä, että koboltin tai koboltin ja jonkin muun metallin seoksen suolaliuos on sulfaatin, nitraatin tai kloridin muodossa.
- 15 4. Patenttivaatimuksen 1 mukainen menetelmä, tunnettu siitä, että suolaliuoksen konsentraatio on välillä 10 ja 120 g/l kokonaismetallipitoisuuden suhteen laskettuna.
 - 5. Patenttivaatimuksen 1 mukainen menetelmä, tunnettu siitä, että kompleksinmuodostaja-aine on ammoniumsulfaattia tai vesipitoista ammoniakkia.
- 6. Patenttivaatimuksen 1 mukainen menetelmä, tunnettu siitä, että kompleksin-20 muodostaja-aineen ja metallin välinen moolisuhde on noin 1,5-2.
 - 7. Patenttivaatimuksen 1 mukainen menetelmä, tunnettu siitä, että pH:n säätämiseen käytetään NaOH:ia.
 - 8. Patenttivaatimuksen 1 mukainen menetelmä, tunnettu siitä, että reaktio suoritetaan lämpötilassa noin 40-90 °C.
- 9. Patenttivaatimuksen 8 mukainen menetelmä, tunnettu siitä, että reaktio suoritetaan lämpötilassa noin 70 °C.
 - 10. Patenttivaatimuksen 1 mukainen menetelmä, tunnettu siitä, että muina metalleina käytetään nikkeliä, mangaania, mangnesiumia tai alumiinia tai näiden seoksia.



- 11. Kobolttihydroksidi tai koboltin ja jonkin muun metallin muodostama seoshydroksidi, **tunnettu** siitä, että sen tilavuuspaino on noin 0,5-2,2 g/cm³, partikkelikoko yli noin 1 μm, tyypillisesti noin 1-20 μm ja omiaispinta-ala noin 0,5-20 m²/g.
- 12. Patenttivaatimuksen 11 mukainen kobolttihydroksidi tai koboltin ja jonkin muun metallin muodostama seoshydroksidi, tunnettu siitä, että muu metalli on nikkeli, mangaani, mangnesium tai alumiini tai näiden seos.
 - 13. Patenttivaatimuksen 11 tai 12 mukainen kobolttihydroksidi tai koboltin ja jonkin muun metallin muodostama seoshydroksidi, tunnettu siitä, että se on valmistettu jollain patenttivaatimusten 1-10 mukaisella menetelmällä.



Keksintö koskee menetelmää korkean tilavuuspainon ja suuren partikkelikoon omaavan kobolttihydroksidin tai koboltin ja jonkin muun metallin seoshydroksidin valmistamiseksi, jossa koboltin vesipitoiseen suolaliuokseen tai koboltin ja jonkin muun metallin seoksen vesipitoiseen suolaliuokseen lisätään alkalisissa olosuhteissa kompleksinmuodostaja-ainetta ja hydroksidi-ionia metallihydroksidin muodostamiseksi. Menetelmässä kompleksinmuodostaja-aine valitaan siten, että se muodostaa ammoniumkompleksin metalli-ionin kanssa. Kompleksinmuodostaja-aineen ja metallin välinen moolisuhde on noin 0,5-3, ja pH säädetään välille 10-13.